

# National Institute of Standards & Technology

# Certificate of Analysis

## Standard Reference Material 3144

### Spectrometric Standard Solution

#### Rhodium

#### Batch Code 490512

This Standard Reference Material (SRM) is intended for use in atomic absorption spectrometry, inductively coupled plasma spectrometry, spectrophotometry, or any other analytical technique that requires aqueous standard solutions for calibrating instruments. SRM 3144 is a single element solution prepared gravimetrically to contain a nominal 1 mg/mL of rhodium with a hydrochloric acid concentration (v/v) of 10 percent. The certified value is based on the gravimetric preparation of the solution and the assay of the rhodium salt. The density of the solution at 22 °C is 1.017 g/mL.

Metal	Concentration (mg/mL)	Source Purity, %	Acid Conc. (v/v) Approximate
Rh	$1.000 \pm 0.003$	(NH <sub>4</sub> ) <sub>3</sub> RhCl <sub>6</sub> *	HCl, 10%

<sup>\*</sup>This material was reduced to rhodium metal which was then analyzed by direct current arc optical emission spectrometry and found to contain 85  $\mu$ g/g Ir, 20  $\mu$ g/g Pt, 70  $\mu$ g/g Ru, 60  $\mu$ g/g Si, and 10  $\mu$ g/g Ag.

The uncertainty in the certified value is calculated as

$$U = 2u_c$$

where  $u_c$  is the "combined uncertainty" calculated according to the ISO Guide [1]. The value  $u_c$  is intended to represent, at the level of one standard deviation, the combined effects of any errors attributable to weighing and uncertainty associated with the dilution as well as those associated with the gravimetric assay of the ammonium hexachlororhodate salt for percent rhodium.

The combined uncertainty consists of Type B components due to uncertainty in the material purity and uncertainty in the material handling, and dilution.

SRM 3144 was prepared by T.A. Butler of the NIST Analytical Chemistry Division. The gravimetric assay procedures were performed by C.M. Beck II of the NIST Analytical Chemistry Division. Direct-current arc spectrometry analyses were performed by C.L. Maul, formerly of Ledoux & Co., Teaneck, NJ, and oxygen and nitrogen analyses were performed by L.W. Ollila and M.P. Ollila of Luvak Inc., Boylston, MA. Technical advice on the gravimetric assay procedures for the rhodium salt was given by P. Blumberg of Ledoux & Co. and E.W. Hobart, Jr., formerly of Ledoux & Co.

Gaithersburg, MD 20899 January 24, 1995

Thomas E. Gills, Chief Standard Reference Materials Program

The support aspects involved in the preparation, certification, and issuance of this SRM were coordinated through the Standard Reference Materials Program by J.S. Kane

#### Procedures for Use

**Stability:** This certification is valid for one year from the shipping date, provided the solutions are kept tightly capped and stored under normal laboratory conditions. NIST will monitor the stability of representative solutions from this SRM lot, and if any changes occur that invalidate this certification, NIST will notify purchasers.

Preparation of Working Standard Solutions: Solutions should be brought to  $22 \pm 1$  °C before use and all glass or plastic surfaces coming into contact with the standard must have been previously cleaned. A working standard solution can be prepared from the SRM solution by serial dilution. Dilutions should be made with certified volumetric class A flasks and 5 or 10 mL class A pipets. All volumetric transfers of solutions should be performed using a proven analytical technique. Each dilution should be acidified with an appropriate high-purity acid and diluted to calibrated volume using high-purity water. The stability of the working standard solution will depend on the final acid concentration; therefore, care should be exercised to ensure that the final acid concentration of the dilution closely approximates that of the SRM. To achieve the highest accuracy, the analyst should prepare daily working solutions from  $100 \mu g/mL$  dilutions of the original SRM solution.

The water-soluble rhodium salt used to prepare this solution standard was gravimetrically assayed for rhodium by first oxidizing the salt to rhodium oxide and then reducing the oxide to rhodium metal with hydrogen in a Rose crucible. Impurities were determined in the resulting rhodium metal and an appropriate deduction made. Independently, high-purity rhodium metal was used to prepare a different water soluble rhodium salt by chlorination of the metal and subsequent reaction with sodium chloride and chlorine in a tube furnace. Inductively-coupled plasma emission spectrometric comparisons of a previous batch of rhodium solutions prepared from each salt were shown to be indistinguishable to 0.2% relative. The details of these procedures are published in reference [2].

#### REFERENCES

- [1] "Guide to the Expression of Uncertainty in Measurement", ISBN 92-67-10188-9, 1st Ed. ISO, Geneva, Switzerland, 1993.
- [2] Beck, C.M. II, Salit, M.L., Watters, R.L. Jr., Butler, T.A. and Wood, L.J. "The Preparation and Certification of a Rhodium Standard Reference Material Solution," *Anal. Chem.*, **1993**, *65*, 2899-2902.